## **AMENDMENTS TO THE CLAIMS:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

## **LISTING OF CLAIMS:**

- 1. (Original) A method for detecting cytochrome c in a given biological sample, comprising: adding to said sample an efficient amount of two redox couples allowing for a cycling oxido-reduction of cytochrome c, said couples comprising an oxidizing agent consisting of cytochrome c oxidase enzyme and a reducing agent specific for cytochrome c with a reduced co-factor; measuring, by a biophysical system depending on the co-factor and allowing to distinguish the co-factor oxidized form from the reduced form, the oxidation of the co-factor which is oxidized during said cycling redox reaction; the amount of the co-factor oxidized form being correlated to the concentration of cytochrome c in the sample.
- 2. (Original) The method of claim 1, wherein said measurement is compared to measurements performed with standard cytochrome c.
- 3. (Currently Amended) The method of claim 1 or 2, wherein the reducing agent is NADH-cytochrome c reductase or NADPH-cytochrome c reductase and the reduced co-factor is NADH or NADPH respectively.
- 4. (Currently Amended) The method of claim 1 to 3, wherein the co-factor is detected by absorption spectrophotometry at 340 nm.

- 5. (Currently Amended) The method of any of claims claim 1 to 4, wherein said agents are, for example but not limited to, under liquid, dried or lyophilised form and obtained by purification of recombinant or natural compounds or by chemical synthesis.
- 6. (Currently Amended) The method of any of claims claim 1 to 5, optimized for any new screening protocol or adaptaded to any existing screening procedure.
- 7. (Original) A kit for detecting cytochrome c in sample to be tested, comprising two redox couples for a cycling oxido-reduction of cytochrome c, said couples comprising an oxidizing agent consisting of cytochrome c oxidase enzyme and a reducing agent specific for cytochrome c with a reduced co-factor.
- 8. (Original) The kit of claim 7, wherein the reducing agent is NADH-cytochrome c reductase and the co-factor is NADH.
- 9. (Original) The kit of claim 7, wherein the reducing agent is NADPH-cytochrome c reductase and the co-factor is NADPH.
- 10. (Currently Amended) The kit of any of claims claim 7 to 9, further comprising cytochrome c as a reference standard.
- 11. (Currently Amended) The kit of any of claims claim 7 to 10, further comprising a buffer.
- 12. (Currently Amended) The kit of claims claim 7 to 11, wherein said agents are, for example but not limited to, under liquid, dried or lyophilised. form, and

obtained by purification of recombinant or natural compounds or by chemical synthesis.

- 13. (Currently Amended) The kit of claims claim 7 to 12, defined for laboratory research only.
- 14. (Currently Amended) The kit of claims claim 7 to 12, defined for diagnostic use.
- 15. (Currently Amended) The kit of any of claims claim 7 to 14, optimized for any format of container, for example but not limited to, 96-well microplates, 384-well microplates, 1 mL cuvettes.
- 16. (Currently Amended) The kit of any of claims claim 7 to 15, optimized for detecting cytochrome c in mitochondrial supernatants.
- 17. (Currently Amended) The kit of any of claims claim 7 to 15, optimized for detecting cytochrome c in cytosol extracts.
- 18. (Currently Amended) The kit of any of claims claim 7 to 15, optimized for detecting cytochrome c in any other biological sample expected to contain cytochrome c.
- 19. (Original) The kit of claim 18, with reagents supplied for the preparation of mitochondrial and/or cytosolic fractions.
- 20. (Original) The kit of claim 19, with methodology for the preparation of mitochondrial and/or cytosolic fractions.